This listing of claims will replace all prior versions, and listings, of claims in the application;

Listing of Claims:

1. (Currently Amended) Compounds of the formula I

$$R^1$$
 R^2
 $C_{mH_{2m}}$
 $C_{mH_{2m}}$
 $C_{mH_{2m}}$
 $C_{mH_{2m}}$
 $C_{mH_{2m}}$
 $C_{mH_{2m}}$

X = N or CH.

 R^1 , R^3 = independently of one another H, OH, OA, CN, Hal, COR⁴ or CH₂R⁴, R^2 = H, an optionally mono- or poly-Hal-substituted, linear or branched alkyl moiety having 1-6 C atoms, or an alkaryl, alkheteroaryl, or heteroaryl moiety. R^4 = OH, OA, NH, NHB or NB₂.

A, B = independently of one another alkyl having 1-6 C atoms,

m = 2, 3, 4, 5 or 6 and

n = 0, 1, 2, 3 or 4,

andor physiologically acceptable salts, derivatives, solvates andor stereoisomers thereof, including mixtures thereof in all ratios.

2. (Original) Compounds according to Claim 1 in which

X = N,

 R^{1} , R^{3} = independently of one another CN, COR^{4} or $CH_{2}R^{4}$,

 R^2 = a linear or branched alkyl having 1-6 C atoms, alkaryl, alkheteroaryl, or heteroaryl,

 $R^4 = OH$, NH_2 , NHB or NB_2 ,

A, B = independently of one another alkyl having 1-6 C atoms,

m = 4 and

n = 0.

and physiologically acceptable salts, derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios.

- 3. (Currently Amended) Compounds according to Claim 1 or 2
- a. 5-{4-[4-(5-cyano-1-methyl-1H-indol-3-yl)butyl]piperazin-1-yl}benzofuran-2carboxamide
- 5-{4-[4-(5-cyano-1-ethyl-1H-indol-3-yl)butyl]piperazin-1-yl}benzofuran-2carboxamide
- c. 5-{4-[4-(5-cyano-1-isopropyl-1H-indol-3-yl)butyl]piperazin-1-yl}benzofuran-2carboxamide
- d. 5-{4-[4-(1-benzyl-5-cyano-1H-indol-3-yl)butyl]piperazin-1-yl}benzofuran-2-carboxamide
- e. 5-{4-[4-(5-cyano-1-propyl-1H-indol-3-yl)butyl]piperazin-1-yl}benzofuran-2-carboxamide
- f. 5-{4-[4-(5-cyano-1-pyridin-2-ylmethyl-1H-indol-3-yl)butyl]piperazin-1-yl}benzofuran-2-carboxamide
- g. 5-{4-[4-(5-cyano-1-phenethyl-1H-indol-3-yl)butyl]piperazin-1-yl}benzofuran-2carboxamide
- (Currently Amended) Process for the preparation of the compounds of the formula I, eharacterised in that comprising
- a) reacting a compound of the formula II, in which R¹ and m have the meanings
 indicated in Claim 1 and Y is a halogen or is an alcohol provided with a protecting
 group known to the person skilled in the art,

is reacted with a compound of the formula III, in which R² has the meanings indicated in Claim 1 and Z represents a leaving group known to the person skilled in the art, such as, for example, p-tosyl, trifluoromethanesulfonyl, methanesulfonyl, benzenesulfonyl, Br, Cl or I

and

b) in that the reacting a compound of the formula IV

obtained in accordance with a) is reacted with a compound of the formula V or a salt thereof, in which $R^3,\,X$ and n have the meanings indicated in Claim 1,

$$\underset{\mathsf{HN}}{\overset{\mathsf{C}_{\mathsf{n}}\mathsf{H}_{2\mathsf{n}}}{\overset{\mathsf{C}_{\mathsf{n}}\mathsf{N}}{\overset{\mathsf{C}_{\mathsf{n}}}{\overset{\mathsf{C}_{\mathsf{n}}}{\overset{\mathsf{C}_{\mathsf{n}}}}{\overset{\mathsf{C}_{\mathsf{n}}}}{\overset{\mathsf{C}_{\mathsf{n}}}}{\overset{\mathsf{C}_{\mathsf{n}}}{\overset{\mathsf{C}_{\mathsf{n}}}}{\overset{\mathsf{C}_{\mathsf{n}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}{\overset{\mathsf{C}}}}{\overset{\mathsf{C}}}}{\overset{\mathsf{C}}}}{\overset{\mathsf{C}}}}}}}$$

in a solvent, optionally with addition of base, at the boiling point of the solvent, or

- in that the converting a base of a compound of the formula I is converted into one of its salts by treatment with an acid.
- 5. (Canceled)
- (Currently Amended) Pharmaceutical composition comprising at least one
 compound according to Claim 1 and/or physiologically acceptable salts, derivatives
 solvates andor stereoisomers thereof, including mixtures thereof in all ratios, and a
 pharmaceutically acceptable carrier.
- (Original) Pharmaceutical composition, according to Claim 6 comprising further excipients and/or adjuvants.
- (Currently Amended) Pharmaceutical composition comprising at least one
 compound according to Claim 1 and/or physiologically acceptable salts, derivatives
 solvates andor stereoisomers thereof, including mixtures thereof in all ratios, and at
 least one further medicament active ingredient.
- 9. (Currently Amended) Process for the preparation of a pharmaceutical composition, eharacterised in that comprising bringing a compound according to Claim 1 and/or one of its physiologically acceptable salts, derivatives, solvates and or stereoisomers, including mixtures thereof in all ratios, is brought into a suitable dosage form together with a solid, liquid or semi-liquid excipient or adjuvant.
- 10. (Canceled)

- 11. (Currently Amended) Use of compounds according to Claim 1 and/or physiologically acceptable salts, derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, for the preparation of a medicament A method for the treatment of diseases associated with the serotonin receptor and/or serotonin recuptake, comprising administering to a host in need thereof an effective amount of a compound according to Claim 1 and/or physiologically acceptable salts, solvates or stereoisomers thereof, including mixtures thereof in all ratios.
- 12. (Currently Amended) Use of compounds according to Claim Land/or-physiologically acceptable salts, derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, for the preparation of a medicament as A method of achieving an anxiolytic, antidepressant, neuroleptic and/or antihypertonic effect and/or for positively influencing obsessive-compulsive disorder (OCD), sleeping disorders, tardive dyskinesia, learning disorders, age-dependent memory disorders, eating disorders, such as bulimia or IBS, and/or sexual dysfunctions, comprising administering to a host in need thereof an effective amount of a compound according to claim 1 and/or physiologically acceptable salts, solvates or stereoisomers thereof, including mixtures thereof in all ratios.
- 13. (Currently Amended) Use of compounds according to Claim 1 and/or physiologically acceptable salts, derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, for the preparation of a medicament A method for the treatment of psychoses, schizophrenia, schizo-affective psychosis, cyclothymia, cpilepsy, cramps, depression (sub-types of severe depression and eyelothymic depression), pathogenic anxiety states (sub-types of panic attacks with or without agoraphobia), superexcitation, hyperactivity, stress illnesses, post-traumatic stress disorders, sleeping disorders, narcolepsy, cyclic manic depression, attention disorders in children and youths, severe developmental disorders, and-disorders of social behaviour with mental retardation, obsessive-compulsive disorders in

the narrower (OCD) and broader sense (OCSD), addiction disorders, disorders in nutrient uptake or eating disorders, for example bulimia, obesity or anorexia nervosa, in particular irritable bowel syndrome (IBS), fibromyalgia, and psychiatric symptoms in senile dementia andor Alzheimer's-type dementia, cognitive impairments (learning and memory disorders), in particular age dependent memory disorders, dementia, tardive dyskinesia, neurodegenerative diseases, such as Parkinson's disease, Alzheimer's disease, Huntington's disease, lathyrism, amyotrophic lateral sclerosis, Lewy bodies dementia, Tourette's syndrome, sexual dysfunctions, premenstrual syndrome, acromegaly, hypogonadism, secondary amenorrhoea, undesired puerperal lactation, extrapyramidal motor disorders, for the treatment of side effects arising in the treatment of extrapyramidal motor disorders with conventional anti-Parkinson's medicaments, and of extrapyramidal symptoms (EPS), tension states, side effects of hypertonia treatment induced by neuroleptics (for example with & methyldopa) or for the prophylaxis, treatment and control of cerebral infarctions-(apoplexia-cerebri), such as strokes and cerebral ischaemia, or for the treatment of pain, in particular chronic pain, migraine, CNS trauma, hypoglycaemia, asthma, glaucoma, cytomegaly and for the treatment of other-degenerative retinal diseases, incontinence, tinnitus, or for the treatment of loss of hearing induced by aminoglycoside antibiotics, comprising administering to a host in need thereof an effective amount of a compound according to claim 1 and/or physiologically acceptable salts, solvates or stereoisomers thereof, including mixtures thereof in all ratios.

14. (Currently Amended) A Set (kit) consisting of separate packs of

- an effective amount of a compound according to Claim 1 and/or physiologically acceptable salts, derivatives, solvates andor stereoisomers thereof, including mixtures thereof in all ratios, and
- b) an effective amount of a further medicament active ingredient.